

We claim:

- Sub B1
1. A poly(ADP-ribose) polymerase (PARP) homolog derived from a human or non-human mammal which has an amino acid sequence which has
 - a) a functional NAD⁺ binding domain and
 - b) no zinc finger sequence motif of the general formula

$$CX_2CX_mHX_2C$$

in which m is an integral value from 28 or 30, and the X radicals are, independently of one another, any amino acid.
 2. A PARP homolog as claimed in claim 1, wherein the functional NAD⁺ binding domain comprises one of the following general sequence motifs:

$$PX_n(S/T)GX_3GKGIYFA,$$

$$(S/T)XGLR(I/V)XPX_n(S/T)GX_3GKGIYFA \text{ or}$$

$$LLWHG(S/T)X_7IL(S/T)XGLR(I/V)XPX_n(S/T)GX_3GKGIYFAX_3SKSAXY$$

in which n is an integral value from 1 to 5, and the X radicals are, independently of one another, any amino acid.
 3. A PARP homolog as claimed in either of the preceding claims, comprising at least another one of the following part-sequence motifs:

$$LX_9NX_2YX_2QLLX(D/E)X_{10/11}WGRVG,$$

$$AX_3FXKX_4KTXNXWX_5FX_3PXX,$$

$$QXL(I/L)X_2IX_9MX_{10}PLGKLX_3QIX_6L,$$

$$FYTIXIPHXXFGX_3PP; \text{ and}$$

$$KX_3LX_2LXDIEXAX_2L,$$

in which the X radicals are, independently of one another, any amino acid.
 4. A PARP homolog as claimed in any of the preceding claims, selected from human PARP homologs, which has the amino acid sequence shown in SEQ ID NO: 2 (human PARP2) or SEQ ID NO: 4 or 6 (human PARP3 type 1 or 2); or murine PARP homologs which

Translation of amended sheet annexed IPER

82

have the amino acid sequence shown in SEQ ID NO:8 (mouse PARP long form) or SEQ ID No:10 (mouse PARP short form).

- Sub A1 Amended
5. A binding partner having specificity for PARP homologs as claimed in any of the preceding claims, selected from
- a) antibodies and fragments thereof,
 - b) protein-like compounds which interact with a part-sequence of the protein, and
 - c) low molecular weight effectors which modulate the catalytic PARP activity or another biological function of a PARP molecule.
6. A nucleic acid comprising
- a) a nucleotide sequence coding for at least one PARP homolog as claimed in any of claims 1 to 4, or the complementary nucleotide sequence thereof;
 - b) a nucleotide sequence which hybridizes with a sequence as specified in a) under stringent conditions; or
 - c) nucleotide sequences which are derived from the nucleotide sequences defined in a) and b) through the degeneracy of the genetic code.
7. A nucleic acid as claimed in claim 6, comprising
- a) nucleotides +3 to +1715 shown in SEQ ID NO:1;
 - b) nucleotides +242 to +1843 shown in SEQ ID NO:3;
 - c) nucleotides +221 to +1843 shown in SEQ ID NO:5;
 - d) nucleotides +112 to +1710 shown in SEQ ID NO:7; or
 - e) nucleotides +1 to +1584 shown in SEQ ID NO:9.
8. An expression cassette comprising, under the genetic control of at least one regulatory nucleotide sequence, at least one nucleotide sequence as claimed in either of claims 6 and 7.
9. A recombinant vector comprising at least one expression cassette as claimed in claim 8.
10. A recombinant microorganism comprising at least one recombinant vector as claimed in claim 9.
11. A transgenic mammal comprising a vector as claimed in claim 9.
- Sub A2

83

- Sub A3
12. A PARP-deficient mammal or PARP-deficient eukaryotic cell, in which functional expression of at least one gene which codes for a PARP homolog as claimed in any of claims 1 to 4 is inhibited.
- 5 13. An in vitro detection method for PARP inhibitors, which comprises
- 10 a) incubating an unsupported or supported polyADP-ribosylatable target with a reaction mixture comprising
- a1) a PARP homolog as claimed in any of claims 1 to 4,
- a2) a PARP activator; and
- 15 a3) a PARP inhibitor or an analyte in which at least one PARP inhibitor is suspected;
- b) carrying out the polyADP ribosylation reaction; and
- c) determining the polyADP ribosylation of the target qualitatively or quantitatively.
- 20 14. A method as claimed in claim 13, wherein the PARP homolog is preincubated with the PARP activator and the PARP inhibitor or an analyte in which at least one PARP inhibitor is suspected, before the polyADP ribosylation reaction is carried out.
- 25 Sub A4 15. A method as claimed in either of claims 13 and 14, wherein the polyADP-ribosylatable target is a histone protein.
16. A method as claimed in any of claims 13 to 15, wherein the
- 30 PARP activator is activated DNA.
17. A method as claimed in any of claims 13 to 16, wherein the polyADP ribosylation reaction is started by adding NAD⁺.
- 35 18. A method as claimed in any of claims 13 to 17, wherein the polyADP ribosylation of the supported target is determined using anti-poly(ADP-ribose) antibodies.
19. A method as claimed in any of claims 13 to 17, wherein the
- 40 unsupported target is labeled with an acceptor fluorophore.
20. A method as claimed in claim 19, wherein the polyADP
- 45 ribosylation of the unsupported target is determined using anti-poly(ADP-ribose) antibody which is labeled with a donor fluorophore which is able to transfer energy to the acceptor fluorophore.

Translation of amended sheet annexed to IPER

84

21. A method as claimed in either of claims 19 and 20, wherein the target is biotinylated histone, and the acceptor fluorophore is coupled thereto via avidin or streptavidin.
- Sub AS 22. A method as claimed in either of claims 20 and 21, wherein the anti-poly(ADP-ribose) antibody carries a europium cryptate as donor fluorophore.
23. An in vitro screening method for binding partners for a PARP molecule, which comprises
- 10 a1) immobilizing at least one PARP homolog as claimed in any of claims 1 to 4 on a support;
- b1) contacting the immobilized PARP homolog with an analyte in which at least one binding partner is suspected; and
- 15 c1) determining, where appropriate after an incubation period, analyte constituents bound to the immobilized PARP homolog;
- or
- 20 a2) immobilizing on a support an analyte which comprises at least one possible binding partner for a PARP molecule;
- b2) contacting the immobilized analyte with at least one PARP homolog as claimed in any of claims 1 to 4 for which a
- 25 binding partner is sought; and
- c2) examining the immobilized analyte, where appropriate after an incubation period, for binding of the PARP homolog.
- 30 24. A method for the qualitative or quantitative determination of nucleic acids encoding a PARP homolog as claimed in any of claims 1 to 4, which comprises
- a) incubating a biological sample with a defined amount of an exogenous nucleic acid as claimed in either of claims
- 35 6 and 7, hybridizing under stringent conditions, determining the hybridizing nucleic acids and, where appropriate, comparing with a standard; or
- b) incubating a biological sample with a pair of
- 40 oligonucleotide primers with specificity for a PARP homolog-encoding nucleic acid, amplifying the nucleic acid, determining the amplification product and, where appropriate, comparing with a standard.
- 45 25. A method for the qualitative or quantitative determination of a PARP homolog as claimed in any of claims 1 to 4, which comprises

85

- 5 a) incubating a biological sample with a binding partner specific for a PARP homolog,
b) detecting the binding partner/PARP complex and, where appropriate,
c) comparing the result with a standard.
26. A method as claimed in claim 25, wherein the binding partner is an antibody or a binding fragment thereof, which carries a detectable label where appropriate.
- 10 27. A method as claimed in any of claims 24 to 26 for diagnosing energy deficit-mediated illnesses.
- 15 28. A method for determining the efficacy of PARP effectors, which comprises
a) incubating a PARP homolog as claimed in any of claims 1 to 4 with an analyte which comprises an effector of a physiological or pathological PARP activity; removing the effector again where appropriate; and
20 b) determining the activity of the PARP homolog, where appropriate after adding substrates or cosubstrates.
29. A gene therapy composition, which comprises in a vehicle acceptable for gene therapy a nucleic acid construct which
25 a) comprises an antisense nucleic acid against a coding nucleic acid as claimed in either of claims 6 and 7; or
b) a ribozyme against a nucleic acid as claimed in either of claims 6 and 7; or
c) codes for a specific PARP inhibitor.
- 30 30. A pharmaceutical composition comprising, in a pharmaceutically acceptable vehicle, at least one PARP protein as claimed in any of claims 1 to 4, at least one PARP binding partner as claimed in claim 5 or at least one coding
35 nucleotide sequence as claimed in claim 6 or 7.
31. The use of low molecular weight PARP binding partners as claimed in claim 5 for the manufacture of a pharmaceutical agent for the diagnosis or therapy of pathological states in
40 the development and/or progress of which at least one PARP protein, or a polypeptide derived therefrom, is involved.
32. The use of low molecular weight PARP binding partners as claimed in claim 5 for the manufacture of a pharmaceutical agent for the diagnosis or therapy of pathological states
45 mediated by an energy deficit.